

**K142816 AbbVie J**Jan 14, 2015  
107 days to decisionK142816 · Product code: **KNT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k142816/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Sep 29, 2014
Decision date	Jan 14, 2015
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>AbbVie, Inc.</b>
Location	North Chicago, IL, US
Contact	Katherine Wortley
510(k) history	6 submissions · 4 cleared · 2014-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142816/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 5, 2026